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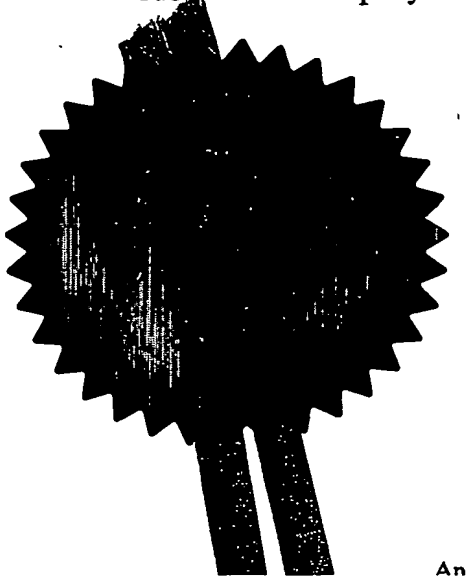

Signed R. Mahoney

Dated 17 October 2003

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Request for grant of a patent

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The Patent Office
Cardiff Road
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Gwent NP10 8QQ

1. Your reference — **4-32680P1**

2. Patent application number *(The Patent Office will fill in this part)* — **0223978.8** — 15 OCT 2002

3. Full name, address and postcode of the or of each applicant *(underline all surnames)*

NOVARTIS AG
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Patent ADP number *(if you know it)* 7125487005

If the applicant is a corporate body, give the country/state of its incorporation — **SWITZERLAND**

4. Title of invention — **Organic compound**

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6. If you are declaring priority from one ore more earlier patent applications, give the country and the date of filing of the or of each of these earlier applications and *(if you know it)* the or each application number

Country	Priority application number *(if you know it)*	Date of filing (day/month/year)

7. If this application is divided or otherwise derived from an earlier UK application, give the number and the filing date of the earlier application

Number of earlier application	Date of filing (day/month/year)

8. Is a statement of inventorship and of right to grant of a patent required in support of this request? *(Answer 'Yes' if:*

a) any applicant named in part 3 is not an inventor, or

b) there is an inventor who is not named as an applicant, or

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(see note (d))

Yes

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Continuation sheets of this form	
Description	13 ✓
Claim(s)	2 ✓
Abstract	
Drawing(s)	

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Priority documents	
Translations of priority documents	
Statement of inventorship and right to grant of a patent *(Patents Form 7/77)*	
Request for preliminary examination and search *(Patents Form 9/77)*	One ✓
Request for substantive examination *(Patents Form 10/77)*	
Any other documents *(please specify)*	

11. I/We request the grant of a patent on the basis of this application

Signature B. A. Yorke & Co. Date 15 October 2002

12. Name and daytime telephone number of person to contact in the United Kingdom

Mrs. E. Cheetham
020 8560 5847

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ORGANIC COMPOUND

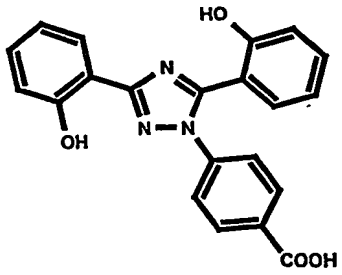
The present invention relates to pharmaceutical dispersible tablets comprising 4-[3,5-bis(2-hydroxyphenyl)-[1,2,4]triazol-1-yl]benzoic acid or a pharmaceutically acceptable salt thereof and is hereinafter referred as Compound I.

Compound I is an orally active iron chelator that is indicated in the treatment of iron overload in transfusion dependent anemias, in particular thalassemia major, thalassemia intermediate and in sickle cell disease to reduce iron-related morbidity and mortality. Compound I can also be used in the treatment of hemochromatosis.

Clinical thalassemia (major and intermedia) are hereditary disorders characterized by defective production of hemoglobin, which leads to decreased production and increased destruction of red blood cells.
Sickle cell disease is caused by a mutation in the hemoglobin-Beta gene leading to the production of abnormal hemoglobin S. Normal red blood cells die after 120 days and sickle cells (red blood cells with hemoglobin S) are destroyed more rapidly (10 to 20 days) causing anemia. This anemia is what gives the disease its commonly known name - sickle cell anemia.
Hemochromatosis, the most common form of iron overload disease, is an inherited disorder that causes the body to absorb and store too much iron. The extra iron builds up in organs and damages them. Without treatment, the disease can cause these organs to fail.

Patients with sickle cell disease or thalassemia, who receive significant numbers of blood transfusions and patients with hemochomatosis require therapy to remove iron from the body, called chelation therapy.

Compound I has the following formula:



Compound I in the free acid form, salts thereof and its crystalline forms are disclosed in the International Patent publication WO97/49395 published on December 31, 1997.

Typically, prescribed daily dosages of Compound I for the treatment of thalassemia are high, e.g. 5 to 40 mg/kg of body weight/day in adults or children. In children, the dosage is preferably 5 to 30 mg/kg of body weight/day. Due to the high dosage strength, the tablet dimensions did not permit to formulate a conventional tablet. Thus, there is a need for an oral dosage form convenient to administer to adults and to children and that provides a daily dosage amount of Compound I. A dispersible tablet is an oral dosage form suitable for high drug loading and pediatric use.

Accordingly, the present invention provides a dispersible tablet with high drug loading comprising Compound I as active ingredient, the active ingredient being present in an amount of from about 5% to 40%, e.g. at least about 10, 15, 20 or 25 %, preferably more than 25% in weight based on the total weight of the dispersible tablet. In particular, the amount of Compound I may vary from 25 to 40%, e.g. 28 to 32% in weight based on the total weight of the dispersible tablet.

The present invention pertains to a dispersible tablet comprising an iron-chelating pharmacologically effective amount of Compound I or a pharmaceutically acceptable salt thereof present in an amount of from 5% to 40% in weight based on the total weight of the tablet.

Compound I may be in the free acid form or pharmaceutically acceptable salts thereof, preferably in the free acid form. The active moiety corresponds to Compound I in the free acid form. Within the context of this disclosure, reference to Compound I is understood to include Compound I in its free acid form or a pharmaceutically acceptable salt thereof or any crystal forms thereof including hydrates or solvates, if not indicated otherwise and where appropriate and expedient.

The present invention also provides a dispersible tablet comprising:
(a) Compound I or a pharmaceutically acceptable salt thereof, and
(b) at least one pharmaceutically acceptable excipient suitable for the preparation of dispersible tablets wherein the amount of Compound I or a pharmaceutically acceptable salt thereof,

calculated as the percentage of the content in weight of the active moiety based on the total the dispersible tablet, is from about 5% to 40% t, e.g. at least about 10, 15, 20 or 25 %, preferably more than 25% in weight based on the total weight of the dispersible table. In particular, the amount of Compound I as active ingredient may vary from 25 to 40%, e.g. 28 to 32% in weight based on the total weight of the dispersible tablet.

In a preferred embodiment of the invention, the present invention provides a dispersible tablet wherein Compound I is in the free acid form (Compound I free acid form).

In a most preferred aspect of the invention, Compound I in the free acid form is in a crystalline form.

One or more pharmaceutically acceptable excipients may be present in the dispersible tablets, e.g. those conventionally used, e.g. (1.1) at least one filler, e.g., lactose, ethylcellulose, microcrystalline cellulose, (1.2) at least one disintegrant, e.g. cross-linked polyvinylpyrrolidinone, e.g. Crospovidone®, (1.3) at least one binder, e.g. polyvinylpyridone, hydroxypropylmethyl cellulose, (1.4) at least one surfactant, e.g. sodium laurylsulfate, (1.5) at least one glidant, e.g. colloidal silicon dioxide, (1.6), at least one lubricant, e.g. magnesium stearate.

Reference is made to the extensive literature on the subject for these and other pharmaceutically acceptable excipients and procedures mentioned herein, see in particular Handbook of Pharmaceutical Excipients, Third Edition, edited by Arthur H. Kibbe, American Pharmaceutical Association, Washington, USA and Pharmaceutical Press, London; and Lexikon der Hilfsstoffe für Pharmazie, Kosmetik and angrenzende Gebiete edited by H.P. Fiedler, 4th Edition, Edito Cantor, Aulendorf and earlier editions which are incorporated herein by reference.

Fillers (1.1) according to the invention are lactose especially lactose monohydrate, preferably lactose monohydrate (200mesh) and lactose spray dried, microcrystalline cellulose especially PH 102, PH 101.

Suitable disintegrants (1.2) according to the invention include but are not restricted to maize starch, CMC-Ca, CMC-Na, microcrystalline cellulose, cross-linked PVP, e.g. as known and

commercially available under the trade names Crospovidone®, Polyplasdone®, available commercially from the ISP company, or Kollidon® XL, alginic acid, sodium alginate and guar gum. Preferably, cross-linked PVP, e.g. Crospovidone® is used.

Binders (1.3) include but are not restricted to starches, e.g. potato, wheat or corn starch, microcrystalline cellulose, e.g. products such as Avicel®, Filtrak®, Heweten® or Pharmacel®; hydroxypropyl cellulose, hydroxyethyl cellulose, hydroxypropylmethyl cellulose, e.g. hydroxypropylmethyl cellulose-Type 2910 USP, hypromellose, and polyvinylpyrrolidone, e.g. Povidone® K30 from BASF. Preferably, polyvinylpyrrolidone is used, most preferably PVP 4K.30.

Appropriate surfactant (1.4) according to the invention is sodium laurylsulfate.

As glidants (1.5), one or more of the following may be used: silica; colloidal silica, e.g. colloidal silica anhydrous, e.g. Aerosil® 200, magnesium trisilicate, powdered cellulose, starch and talc. Preferably, colloidal silicon dioxide is used.

As lubricants (1.6) one or more of the following may be used Mg-, Al- or Ca-stearate, PEG 4000 – 8000, talc, sodium benzoate, glyceryl mono fatty acid, e.g. having a molecular weight of from 200 to 800 Daltons e.g. gylceryl monostearate (e.g., Danisco, UK), glyceryl dibehenate (e.g., CompritolATO888™, Gattefossé France), glyceryl palmito-stearic ester (e.g. Precirol™, Gattefossé France), polyoxyethylene glycol (PEG, BASF), hydrogenated cotton seed oil (Lubitrab, Edward Mendell Co Inc), castor seed oil (Cutina HR, Henkel). Preferably, magnesium stearate is used.

One or more of these pharmaceutically acceptable excipients can be selected and used having regard to the particular desired properties of the dispersible tablet by routine experimentation.

According to the present invention, the amount of filler (1.1) may vary within a range of from about 35 to 55%, in particular 40 to 50% in weight based on the total weight of the dispersible tablet.
The amount of disintegrant (1.2) may vary within a range of from to 5 to 40%, e.g. 10 to 35% in weight based on the total weight of the dispersible tablet.

The amount of binder (1.3) may vary from 1 to 10 %, preferably from 1.5 to 5 % in weight based on the total weight of the dispersible tablet.
The amount of surfactant (1.4) may vary from 0.1 to 2%, preferably from 0.2 to 1%.
The amount of glidant (1.5) may vary within ranges of from 0.1 to 5%, in particular 0.1 to 2.5%, e.g. 0.1 to 0.5% in weight based on the total weight of the dispersible tablet.
The amount of lubricant (1.6) may be below 1% in weight based on the total weight of the dispersible tablet, preferably below 0.5%, most preferably below 0.4% and even most preferably the amount of lubricant is ranging between 0.01% and 0.4%. Very preferably the amount of lubricant is above 0.02% and below 0.4% in weight based on the total weight of the dispersible tablet.

It will be appreciated that any given excipient may serve more than one function e.g. as filler, disintegrant, binder, glidant, and/or lubricant.

In a preferred aspect of the invention, the dispersible tablet comprises the following pharmaceutically acceptable excipients: one or more fillers in a total amount of about 40% to 50% in weight based on the total weight of the dispersible tablet, one or more binders in a total amount of about 1.5% to 5% in weight based on the total weight of the dispersible tablet, one or more disintegrants in a total amount of about 10% to 35% in weight based on the total weight of the dispersible tablet, one or more glidants in a total amount of about 0.1% to 0.5% in weight based on the total weight of the dispersible tablet, and/or one or more lubricants in a total amount of about 0.01 % to 0.4 % in weight based on the total weight of the dispersible tablet.

The absolute amounts of each pharmaceutically acceptable excipient and the amounts relative to other pharmaceutically acceptable excipients is similarly dependent on the desired properties of the dispersible tablet and may also be chosen by routine experimentation.

Procedures which may be used may be conventional or known in the art or based on such procedures e.g. those described in L. Lachman et al. The Theory and Practice of Industrial Pharmacy, 3rd Ed, 1986, H. Sucker et al, Pharmazeutische Technologie, Thieme, 1991, Hagers Handbuch der pharmazeutischen Praxis, 4th Ed. (Springer Verlag, 1971) and Remington's Pharmaceutical Sciences, 13th Ed., (Mack Publ., Co., 1970) or later editions.

The present invention provides dispersible tablets having a disintegration time below 5 minutes. The dispersible tablets of the invention are, despite the high drug loading, dispersible in less than 5 minutes, preferably less than 3 minutes, and, therefore, convenient to administer. This leads to a better patient compliance.

In another embodiment this invention provides a dispersible tablet comprising from 100 mg to 800 mg of Compound I as active ingredient, e.g. of from 100 mg to about 600 mg. Most preferably, dispersible tablets according to the invention are dispersible tablets containing 125 mg, 250 mg or 500 mg of Compound I as active ingredient.

Accordingly, the present invention provides for dispersible tablets, e.g. dispersible tablets, containing an amount of Compound I, equal to 125 mg, 250 mg, or 500 mg of Compound I free acid form. Most preferably, the Compound I in the free acid form used for the dispersible tablet according to the invention is the crystalline form, especially the crystalline form the preparation of which is described in example 5 of WO97/49395.

According to the invention, the process for the preparation of the dispersible tablets consists of granulating an inner phase, mixing it together with one or more pharmaceutically acceptable excipients and compressing the obtained mixture under spray lubrication conditions.

The inner phase comprises Compound I. Preferably the inner phase comprises Compound I and one or more pharmaceutically acceptable excipients. Preferably, the pharmaceutically acceptable excipients of the inner phase are one or more fillers, one or more disintegrants, one or more binders and one or more surfactants. Preferably the amount of one or more fillers in the inner phase is ranging from about 5 to 35% in weight based on the total weight of the dispersible tablet, more preferably 10 to 30% and most preferably 15 to 25%. The filler according to the invention is preferably lactose monohydrate. The disintegrant is preferably Crospovidone XL. The amount of disintegrant present in the inner phase is preferably ranging from 5 to 30%, more preferably 7 to 25% in weight based on the total weight of the dispersible tablet. The Compound I and one or more fillers and one or more disintegrants are mixed together with a wetting solution comprising one or more surfactants, water and one or more binders. The preferred binder is PVP 4K.30. The mixture is processed for granulation, e.g. using a wet high-shear granulator to form the wet-granulates. The wet-granulates may then be, dried, e.g. using a fluid bed dryer and calibrated, e.g., using an oscillating granulator.

The outer phase consists of one or more pharmaceutically acceptable excipients and is mixed with the inner phase using, e.g. a free fall mixer. Preferably, one or more fillers and one or more glidants are added. Most preferably, cellulose microcrystalline and lactose are added as fillers. Even more preferably, microcrystalline cellulose is added in the range of 5 to 20% in weight based on the total weight of the dispersible tablet and lactose is added in the range of 5 to 20% in weight based on the total weight of the dispersible tablet. The outer phase according to the invention may also contain one or more glidants, most preferably colloidal silicon dioxide. In a preferred embodiment, the amount of glidant in the outer phase is ranging from about 0.1 to 5%, preferably 0.1 to 2.5%, most preferably 0.1 to 0.5% in weight based on the total weight of the tablet.

It is a particular aspect of the invention that one or more lubricants, instead of being incorporated into the mixture of the inner and outer phase, are deposited on the punches of the tabletting machine before compression. According to the invention, one or more lubricants are sprayed on the material contacting surfaces of pressing tools (punches) of the tabletting machine before compression.

In one embodiment of the invention, the process for the preparation of a dispersible tablet comprises

(a) forming an inner phase comprising

(i) mixing the Compound I together with pharmaceutically acceptable pharmaceutically acceptable excipients,

(ii) wet-granulating

(b) forming an outer phase comprising

(iii) adding further pharmaceutically acceptable excipients to the inner phase and mixing;

(c) forming the dispersible tablet by

(iv) compressing the mixture obtained in step (iii) under spray lubrication condition.

More specifically, in one aspect the present invention provides a process comprising:

(i) mixing the Compound I and pharmaceutically acceptable excipients, e.g. one or more fillers, e.g. lactose and one or more disintegrants, e.g., Crospovidone XL in a high shear mixer;

(ii) adding a solution of one or more surfactant and one or more binder, subjecting the mixture to wetting/kneading, e.g. in a high shear mixer, wet-granulating using, e.g. a rotating impeller, drying, e.g., in a fluidized bed dryer then calibrating in an oscillating granulator, and;

(iii) adding pharmaceutically acceptable excipients, e.g. sieved excipients, such as one or more fillers, e.g., microcrystalline cellulose, lactose, one or more glidant, e.g. colloidal silicon dioxide, and mixing, e.g. in a free fall mixer;

(iv) tabletting the mixture obtained in step (iii) by compression, e.g. in a conventional tablet press, preferably a rotary machine and spraying the lubricant on the materials contacting surfaces of pressing tools.

By "inner phase" is meant the granulate phase (steps (i) and (ii)) including the active ingredient Compound I and one or more the pharmaceutically acceptable excipients.

By "outer phase" is meant one or more pharmaceutically acceptable excipients added to the inner phase (granulates) (step (iii).

By "total weight of the dispersible tablet" is meant the weight of a tablet being the inner and the outer phase.

The physical and chemical stability may be tested in conventional manner, e.g. the dispersible tablets may be tested as such by measurement of dissolution, friability, disintegration time, assay for Compound I degradation products, appearance and/or microscopy, e.g. after storage at room temperature, i.e. 25°C, and/or storage at 40°C.

The dispersible tablets may vary in shape and be, for example, round, oval, oblong, cylindrical or any other suitable shape. A characteristic of the dispersible tablets according to the invention is their small amount of magnesium stearate having regard to the amount of Compound I contained therein, thus allowing a disintegration time, which complies with the European Pharmacopea Specifications.

In a preferred embodiment of the invention dispersible tablets obtained by the compression method described above are round or oval. The edges of the dispersible tablets may be beveled

or rounded. Most preferably, the dispersible tablets are round with bevelled edges. The dispersible tablets according to the invention may be scored.

The dispersible tablet according to the invention is preferably round, flat with bevelled edges. The 125 mg dispersible tablet has a diameter ranging between 10 and 20 mm, most preferably between 10 and 15 mm. The preferred diameter of the 125 mg dispersible tablet is 12 mm. Its thickness is ranging from 2.5 to 4.5 mm, preferably between 3.2 and 3.9 mm. The 250 mg dispersible tablet has a diameter ranging from 12 to 20 mm, preferably between 14 and 18 mm, the most preferred diameter is 15 mm. Its thickness is ranging from 3.5 to 5.5 mm, most preferably between 4 and 5 mm. The 500 mg dispersible tablet has a diameter ranging from 15 to 30 mm, preferably between 15 and 25 mm, the most preferred diameter is 20 mm. Its thickness is ranging from 4.5 to 6.5 mm, most preferably between 5 and 6 mm.

The dispersible tablets of the invention comprising about 125 mg of Compound I as active moiety may furthermore have a hardness of from about 50 to 120 N, preferably 60 to 100 N. The dispersible tablets of the invention comprising about 250 mg of Compound I may have a hardness of 70 to 150 N, preferably 90 to 130 N. The dispersible tablets of the invention comprising about 500 mg of Compound I may have a hardness of 80 to 190 N, preferably 110 to 160 N.

Preferably, the disintegration time is not more than 5 minutes, most preferably the disintegration time is less than 3 minutes as measured using a desintegration time apparatus.

By "disintegration time" is meant the time that needs the dispersible tablet to disintegrate in water at room temperature in a disintegration time device.

The dispersible tablet of the present invention is dispersible in an aqueous phase, preferably water.

The dispersible tablets of the invention may furthermore be colored and/or marked so as to impart an individual appearance and to make them instantly recognizable. The use of dyes can serve to enhance the appearance as well as to identify the dispersible tablets. Dyes suitable for use in pharmacy typically include carotinoids, iron oxides or chlorophyll. The dispersible tablets of the invention may be marked using an imprint code.

The dispersible tablets of the invention are useful for the treatment of iron overload in transfusion dependent anemias, in particular thalassemia major, thalassemia intermediate and sickle cell disease and in the treatment of hemochromatosis.

The activity and characteristics of the dispersible tablets of the invention may be indicated in standard clinical trials and/or animal trials.

Furthermore, the dispersible tablets of the invention obtained are stable both to the production process and during storage, e.g. for 2 years or even 3 years in conventional packaging, e.g., sealed aluminium blister packs. Less than about 5%, e.g. 2 or 3% or less of Compound I as active ingredient may degrade during this time as determined in conventional tests. For example, less than 1% of Compound I as active ingredient is degraded in one year in HDPE filled bottles.

Depending on age, individual condition, mode of administration, and the clinical picture in question, effective doses, for example daily dosing of dispersible tablets of the invention, comprising, e.g., 350 to 2800 mg are administered to patients of 70 kg body weight.

The invention relates also to a method of administering to a mammal, preferably a human subject in need of such a treatment, Compound I in the form of a dispersible tablet, preferably a dispersible tablet. The invention relates especially to such method wherein a daily dose of 5 to 40 mg/kg of body weight/day of Compound I as active ingredient is administered to a patient. It will be understood that the specific dose level for any particular patient will depend upon a variety of factors including the age, the body weight, general health, drug combination with one or more active drugs, type and severity of the disease.

The medicament package comprises dispersible tablets according to the invention and printed instructions directing that one or more dispersible tablets of Compound I be administered orally.

The following non-limitative examples illustrate the invention.

Example 1: Dispersible tablet Formulation (125 mg, 250 mg and 500 mg dispersible tablets) with a disintegration time above 3 minutes.

Components		%	Amount per dispersible tablet (mg)		
			125 mg	250 mg	500 mg
Phase I	Compound I (free acid form)	29.41	125.0	250.00	500.0
	Lactose 200 Mesh (1.1)	22.09	93.88	187.75	375.5
	Crospovidone XL (1.2)	10.00	42.50	85.00	170.00
Phase II	PVP 4K.30 (1.3)	3.00	12.75	25.50	51.00
	Sodium laurylsulfate (1.4)	0.50	2.13	4.25	8.50
Phase III	Crospovidone XL (1.2)	10.00	42.50	85.00	170.00
	Microcrystalline cellulose (1.1)	11.90	50.57	101.15	202.3
	Lactose spray dried (1.1)	11.90	50.57	101.15	202.3
	Aerosil 200 (1.5)	0.20	0.85	1.70	3.40
Phase IV	Magnesium stearate (1.6)	1.00	4.25	8.50	17.00
Tablet weight (mg)		100.00	425.00	850.00	1700.00
Tablet diameter (mm)		-	12	15	20

Example 2: Dispersible tablet Formulation (125 mg, 250 mg and 500 mg dispersible tablets) with a disintegration time below 3 minutes.

Components		%	Amount per dispersible tablet (mg)		
			125 mg	250 mg	500 mg
Phase I	Compound I (free acid form)	29.4	125.0	250.0	500.0
	Lactose 200 Mesh (1.1)	17.1	72.6	145.2	290.4
	Crospovidone XL (1.2)	15.0	63.7	127.4	254.8
Phase II	PVP 4K.30 (1.3)	3.0	12.8	25.6	51.2
	Sodium laurylsulfate (1.4)	0.5	2.1	4.2	8.4
Phase III	Crospovidone XL (1.2)	5.0	21.3	42.6	85.2
	Microcrystalline cellulose (1.1)	14.9	63.3	126.6	253.2

	Lactose spray dried (1.1)	14.9	63.3	126.6	253.2
	Aerosil 200 (1.5)	0.2	0.9	1.8	3.6
Phase IV	Magnesium stearate (1.6)	< 0.2*			
Tablet weight (mg)		100.0	425	850	1700
Tablet diameter (mm)		-	12	15	20
Tablet thickness (mm)		-	3.6+/-0.3	4.5+/-0.3	5.5+/-0.3

Dispersible tablets of Compound I free acid according to the invention were prepared by forming a inner phase by wet granulation of a mixture of Phase I and Phase II ingredients, Phase III ingredients formed the outer phase and the lubricant (Phase IV) is sprayed directly onto the punches of the tabletting machine.

Example 3: Properties of the 125 mg dispersible tablet of Example 2

Test	release specifications
Tablet shape	12 mm diameter, round, flat, bevelled edge with engraving (IA/NVR)
Tablet appearance	Off white
Friability	Max. 1% (0 unit broken)
Crushing strength (mean)	Mean >=70N
Dissolution rate	Q =75 % within 30 min
Disintegration time	All units < 3 min
Average mass	403.75 – 446.25 mg
Mean content	95.0 –105.0 %

Example 4: Properties of the 250 mg dispersible tablet of Example 2

Test	release specifications
Tablet shape	15 mm diameter, round, flat, bevelled edge with engraving (IB/NVR)
Tablet appearance	Off white
Friability	Max. 1% (0 unit broken)
Crushing strength (mean)	Mean >=90N
Dissolution rate	Q =75 % within 30 min
Disintegration time	All units < 3 min
Average mass	807.5 – 992.5 mg
Mean content	95.0 –105.0 %

Example 5: Properties of the 500 mg dispersible tablet of Example 2

Test	release specifications
Tablet shape	20 mm diameter, round, flat, bevelled edge with engraving (IC/NVR)
Tablet appearance	Off white
Friability	Max. 1% (0 unit broken)
Crushing strength (mean)	Mean >=110N
Dissolution rate	Q =70 % within 30 min
Disintegration time	All units < 3 min
Average mass	1615 – 1785 mg
Mean content	95.0 –105.0 %

Example 6: Magnesium Stearate Assay

	125 mg tablet			250 mg tablet		500 mg tablet		
	L.P.	Pilot	FSP	P.T.	FSP	L.P.	Pilot	FSP
Min (%w/w)	0.1	0.09	0.04	0.08	N/A	0.04	0.03	0.04
Max (%w/w)	0.24	0.36	0.14	0.16	N/A	0.10	0.14	0.06

L.P.: laboratory phase, Pilot: Pilot phase (x 2 of the size of the batch of the laboratory phase), FSP: full scale pilot (batches of production size), P.T.: preliminary trial, N/A: results not yet available; 0.1% w/w of magnesium stearate is equivalent to 1000 ppm.

CLAIMS

1. A dispersible tablet comprising Compound I or a pharmaceutically acceptable salt thereof present in an amount of from 5% to 40% in weight based on the total weight of the tablet.

2. A dispersible tablet comprising (a) Compound I or a pharmaceutically acceptable salt thereof, and (b) at least one pharmaceutically acceptable excipient suitable for the preparation of tablets, wherein the Compound I or a pharmaceutically acceptable salt thereof is present in an amount of from 5% to 40% in weight based on the total weight of the tablet.

3. A dispersible tablet comprising an iron-chelating pharmacologically effective amount of Compound I or a pharmaceutically acceptable salt thereof present in an amount of from 5% to 40% in weight based on the total weight of the tablet.

4. The dispersible tablet according to claim 1, 2 or 3 wherein the Compound I is in the free acid form.

5. The dispersible tablet according to any one of claims 1 to 4 wherein Compound I is in a crystalline form.

6. The dispersible tablet according to any one of claims 1 to 5 wherein a lubricant is present in less than 1% in weight based on the total weight of the tablet.

7. The dispersible tablet according to claim 6 wherein the lubricant is present in less than 0.4% in weight based on the total weight of the tablet.

8. The dispersible tablet according to any one of claims 1 to 7 wherein the disintegration time of the tablet is of 5 minutes or less.

9. The dispersible tablet according to any one of claims 1 to 7 wherein the disintegration time of the tablet is of 3 minutes or less.

10. The dispersible tablet according to any one of claims 1 to 9 wherein the pharmaceutically acceptable excipients comprise:
(i) at least one filler in a total amount of 35 to 55 % in weight based on the total weight of the tablet,
(ii) at least one disintegrant in a total amount of about 10% to 35% in weight based on the total weight of the tablet
(iii) at least one binder in a total amount of about 1.5% to 5% in weight based on the total weight of the tablet,
(iv) at least one surfactant in a total amount of about 0.2% to 1% in weight based on the total weight of the tablet,
(v) at least one glidant in a total amount of about 0.1% to 0.5% in weight based on the total weight of the tablet, and/or
(vi) at least one lubricant in a total amount of less than 0.4% in weight based on the total weight of the tablet.

11. The dispersible tablet according to any one of claims 1 to 10 wherein the lubricant is magnesium stearate.

12. The dispersible tablet according to anyone of claims 1 to 11 containing an amount of Compound I of about 100 mg to 600 mg of Compound I in its free acid form.

13. A method of administering to a mammal in need of such a treatment a daily dose of 5 to 40 mg/kg of body weight/day of Compound I as active ingredient.

14. A process for the preparation of the dispersible tablet according to any one of the preceding claims, which process comprises
(i) mixing the Compound I or a pharmaceutically acceptable salt thereof and at least one pharmaceutically acceptable excipient;
(ii) wet-granulating;
(iii) mixing with at least one pharmaceutically acceptable excipient to form a mixture; and
(iv) spraying the lubricant on the materials contacting surfaces of pressing tools of the tabletting machine and compressing the mixture obtained in step (iii) to form a tablet.

15. The process according to claim 14 wherein the lubricant is magnesium stearate.

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